

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

**As rescanning documents *will not* correct images,
please do not report the images to the
Image Problem Mailbox.**

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶: A61K 9/70, 31/485	A1	(11) International Publication Number: WO 97/39742 (43) International Publication Date: 30 October 1997 (30.10.97)
(21) International Application Number: PCT/SE97/00484 (22) International Filing Date: 21 March 1997 (21.03.97) (30) Priority Data: 9601528-4 23 April 1996 (23.04.96) SE (71) Applicant (for all designated States except US): PHARMACIA & UPJOHN AB [SE/SE]; S-112 87 Stockholm (SE). (72) Inventors; and (75) Inventors/Applicants (for US only): HOECK, Ulla [DK/DK]; Kighusvaenget 7, DK-3400 Hillerød (DK). KREILGÅRD, Bo [DK/DK]; Smedievej 18, DK-3400 Hillerød (DK). KRISTENSEN, Helle [DK/DK]; Lindegårds Allé 16, DK-3550 Slangerup (DK). (74) Agent: HEDENSTRÖM, John; Pharmacia & Upjohn AB, P.O. Box 941, S-251 09 Helsingborg (SE).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: TRANSDERMALLY ADMINISTERED DEXTROMETHORPHAN AS ANTITUSSIVE AGENT		
(57) Abstract Device for transdermal administration of dextromethorphan, (+)-3-methoxy-17-methyl-9a,13a,14a-morphinan, optionally encompassing salts, prodrugs and metabolites thereof, optionally together with pharmaceutically acceptable carrier(s) to a human being or an animal in order to achieve an antitussive effect. Use of an antitussive compound comprising dextromethorphan, (+)-3-methoxy-17-methyl-9a,13a,14a-morphinan, optionally encompassing salts, prodrugs and metabolites thereof, and optionally together with pharmaceutically acceptable carrier(s), for the manufacture of a composition to be administered transdermally for achieving an antitussive effect. Method for achieving an antitussive effect in a living body by transdermal administration of a compound comprising dextromethorphan, (+)-3-methoxy-17-methyl-9a,13a,14a-morphinan, optionally encompassing salts, prodrugs and metabolites thereof, and optionally together with pharmaceutically acceptable carrier(s).		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

TRANSDERMALLY ADMINISTERED DEXTROMETHORPHAN AS ANTITUSSIVE AGENT

Field of invention

This invention relates to use of dextromethorphan, optionally encompassing salts, prodrugs and metabolites thereof, for the manufacturing of a medicament to be administered transdermally for achieving an antitussive effect and to methods of treating diseases being treatable with antitussive agents by transdermal administration of dextromethorphan, optionally encompassing salts, prodrugs and metabolites thereof.

10 Background

Dextromethorphan, (+)-3-methoxy-17-methyl-9a,13a,14a-morphinan, is a synthetic opioid. Normally the hydrobromide of dextromethorphan is used pharmacologically, although other salts are not excluded. The preparation of (+)-3-methoxy-17-methyl-9a,13a,14a-morphinan was disclosed in US patent 2,676,177 (SCHNIDER ET AL.) and in Häfliger et al., *Helv. Chim. Acta* 39, 1956: 2053.

Clinically, in connection with tussometri dextromethorphan has shown a significant effect on reducing coughing frequency as well as intensity compared to placebo at a dosage of 40 mg perorally, an effect of the same order of magnitude as 60 mg codeine, see Mathys, *Schweiz Med Wschr* 1985;115: 307-11. However dextromethorphan has not shown any antitussive effect upon inhalation of 1 - 30 mg. Also demethylated metabolites, including dextrorphan, have shown cough suppressing effects, see Martindale, *The Pharmaceutical Press*, London, 1993: 746.

25 Dextromethorphan is a safe drug as concluded by Bem J.L., Peck R., *Drug safety*, 1992 (7): 190-199. Dextromethorphan has fewer side-effects than the other antitussive agents codeine and noscapine.

Dextromethorphan is rapidly converted in the liver into inter alia dextrorphan, which also has a clinical activity, see above, however on other receptors than dextromethorphan. Pharmacokinetic studies have shown that populations can be divided into two main groups based on their ability to

metabolize dextromethorphan, the so called poor metabolizers and the extensive metabolizers, see e.g. J.-C. Duché et al., "Dextromethorphan O-demethylation and dextrorphan glucoronidation in a French population", *Int J. of Clin Pharm, Therapy and Tox*, 1993; 31(8):392-98, J.S. Marinac et al.,
5 "Dextromethorphan Polymorphic Hepatic Oxidation (CYP2D6) in Healthy Black American Adult Subjects", *Therapeutic Drug Monitoring*, Raven Press New York, 1995; 17:120-124, and Chen et al., "Dextromethorphan: pharmacogenetics, and a pilot study to determine its disposition and antitussive effect in poor and extensive metabolisers", *Eur. J. Pharmacol* 1990;
10 183(4):1573-74. Around 10 % of the population are slow metabolizers of dextromethorphan and therefore more easily have side-effects, most often being fairly mild, such as drowsiness, confused speech, nausea and dizziness, although serious in case of overdosing, such as excitation, confusion and respiratory depression. The clinical implications of these findings are that
15 different dosing regimes should be used for the individual patients. As this difference is related to the first-pass metabolism in the liver it is highly advantageous to avoid the first pass passage of the drug.

As metabolism following transdermal delivery of a drug is of much lesser extent than after oral delivery of the drug it is highly desirable to deliver dextromethorphan through the transdermal route.
20

When administered perorally dextromethorphan undergoes an extensive first-pass metabolism, i.e. the oral bioavailability is low meaning that fairly high doses need to be given. Absolute bioavailabilities have been reported as low as 3.8 % in dogs, see Barnhart J.W., Massad E.N.,
25 "Determination of dextromethorphan in serum by gas chromatography", *J. Chromatography* 1979, 163: 390-395. Other reported values are 7 % and 18 %, see Dixon et al., *Res. Commun. Chem. Pathol. Pharmacol.*, 1978;22:243).

The half-life of dextromethorphan is around 4 - 6 hours, which means
30 that the plasma concentration varies substantially during day and night unless dextromethorphan is delivered frequently, by peroral administration at

least 3 - 4 times daily. Even then the sleeping pattern of the patient will be disturbed by cough attacks as the antitussive effect will not remain through a whole night. The sleeping pattern disturbance, as well as the other adverse effects mentioned above, are removed or reduced with the present invention being transdermally administered dextromethorphan as antitussive agent. The above transdermal administration can be used for human beings as well as animals.

Prior Art

Transdermal administration of dextromethorphan, but not as anti-tussive agent, is known, e.g. from US 5,260,066 (CARLTON ET AL.) for cryogel bandages. Here dextromethorphan is administered only to sites of trauma, column 2, lines 59-60, whereas in the present invention dextromethorphan is only administered to intact skin. Further US 5,260,066 just mentions dextromethorphan in a long listing of drugs. There are no examples showing administration of dextromethorphan. Further US 5,260,066 does not even mention that an antitussive effect should be achieved. Supposedly this is not what is desired upon administration to sites of trauma.

WO 91/15261 (MEDTRONIC) concerns iontophoretic devices which depend upon the physical activity of the patient and just mentions dextromethorphan on page 4, line 32-33, as a drug which could possibly be administered via said devices. But there are no examples showing that this is at all possible with said devices. Dextromethorphan is further not mentioned in the claims. Thus, WO 91/15261 simply concerns a very special device, requiring measurement of patient activity (page 4, lines 24-25, which means an activity sensor (page 11, lines 10-17). This is a non-useful device for administering dextromethorphan as the administration takes place once the patient starts coughing - which is too late. Thus, WO 91/15261 is in all respects an irrelevant and non-enabling reference. WO 91/15261 corresponds to US 5,213,568 (LATTIN ET AL.) which thus also is a non-relevant reference.

WO 95/05416 (CYGNUS THERAPEUTIC SYSTEMS) discloses mucoadhesive devices for administration of drugs, inter alia dextromethorphan, to a body cavity, specifically to the oral cavity. It does not relate to transdermal administration.

5 US 4,783,450 (FAWZI ET AL.), corresponding to WO 88/07871 (WARNER LAMBERT) discloses the use of lecithin for enhancing transdermal penetration. US 4,645,502 (GRACE ET AL.) discloses a specific system for transdermal delivery of highly ionized fat insoluble drugs. WO 93/07902 (RICHARDSON-VICKS, INC.) discloses compositions for topical application comprising a drug
10 and a non-ionic polyacrylamide. WO 93/07903 (RICHARDSON-VICKS, INC.) discloses compositions for topical application comprising a drug and a high molecular weight cationic polymer. EP 0 351 897 (THE PROCTER & GAMBLE COMPANY) discloses pharmaceutical compositions comprising a drug, a fatty acid and an alkane diol. EP 0 349 763 (BRISTOL-MYERS COMPANY) discloses
15 a composition for trans-dermal administration comprising a drug and an imidazole derivative as penetration enhancer. US 4,888,354 (CHANG ET AL.) discloses compositions for topical administration of drugs present in both free and acid addition salt form. US 4,557,934 (COOPER) discloses topical compositions comprising a drug and 1-dodecyl-azacycloheptan-2-one as
20 penetration enhancing agent. In all these patent documents dextromethorphan is just mentioned in lengthy listings of drugs which theoretically might be included in the claimed compositions.

Anyhow there are nowhere in the above patent documents any examples of formulations including dextromethorphan as an antitussive agent.
25 Thus use of transdermally administered dextromethorphan as an antitussive agent has neither been contemplated nor shown.

The only non-patent literature reference relating to transdermal delivery of dextromethorphan being known to the applicant is Mahjour et al., J. Controlled Release 14 (3); 1990:243-252. The contents thereof essentially
30 corresponds to the above mentioned patent US 4,783,450 (FAWZI ET AL.).

Hence the present invention, as further described below, is both new and inventive over prior art.

Objects of the invention

Disturbance of sleeping pattern and the other above mentioned disadvantages are removed or reduced when dextromethorphan is administered transdermally.

Accordingly, a first object of the present invention is to provide a device for transdermal administration use of dextromethorphan, optionally encompassing salts, prodrugs and metabolites thereof, for achieving an antitussive effect. The administration can be to a human being or to an animal. The antitussive effect is for treating, including suppressing, any kind of irritant cough, such as, but not exclusively, non-productive and dry coughs.

A second object of the invention is to provide use of an antitussive compound comprising dextromethorphan for the manufacture of a composition to be administered transdermally for treating cough or conditions associated with cough.

A third object of the invention is to provide a method of treating diseases, in humans or animals, which are treatable with antitussive agents by administering dextromethorphan transdermally.

Other objects of the invention will become apparent to one skilled in the art, and still other objects will become apparent hereinafter.

Summary of the invention

The present invention relates to transdermal administration of dextromethorphan, optionally encompassing salts, prodrugs and metabolites thereof for achieving an antitussive effect. This effect is primarily achieved through the systemic effect of dextromethorphan. Anyhow other actions are not excluded.

Brief description of the drawings

Figures 1A - 1D are schematic drawings of different types of devices for transdermal delivery of drugs.

Figure 2 is a diagram showing *in vitro* skin permeation of dextromethorphan from different solvents according to Example 3.

Figure 3 is a diagram showing *in vitro* permeation of dextromethorphan through different membranes in accordance with Example 4.

5 Figure 4 is a diagram showing *in vitro* release of dextromethorphan from different transdermal systems in accordance with Examples 5 and 6.

Detailed description of the invention

Transdermal delivery of drugs can be achieved from topical products such as ointments or cremes or from transdermal devices. The present invention relates to administration via transdermal devices, which usually are
10 called transdermal patches.

Devices usable as transdermal patches can be categorized in many different ways. A comprehensive categorization of transdermal devices is found in Steven Wick, "Developing A Drug-In-Adhesive Design For Transdermal
15 Drug Delivery", Adhesives Age, 1995; 38(10):18 - 24, which hereby is incorporated by reference. Wick essentially divides transdermal devices into the below four main groups:

- the reservoir type, in which the drug is placed in a liquid or a gel and delivered to the skin across a rate-moderating membrane;
- 20 - the matrix type, in which the drug is placed within a non-adhesive polymeric material, typically a hydrogel or soft polymer;
- the drug-in-adhesive type, in which the drug is placed within an adhesive polymer;
- the multi-laminate type, which is similar to the drug-in-adhesive design but which incorporates an additional layer of pressure sensitive adhesive to cover the entire device and affix it to the skin.
25

The above four main types of transdermal devices are schematically illustrated in FIG. 1A - 1D.

A fifth important type, not mentioned by Wick, is the iontophoretic
30 type, in which an electrical potential gradient is used for transferring the drug through the skin - see further e.g. Parminder Singh et al, "Iontophoresis

in Drug Delivery: Basic Principles and Applications", Critical Reviews in Therapeutic Drug Carrier Systems, 1994; 11 (2&3):161-213.

The above split-up into groups is not very strict as variations and combinations of each may be envisaged. So may a multi-laminate type device encompass a device with many layers in a sandwich construction, such as the drug in one layer, excipients such as enhancers in a further layer, a membrane in another layer and an adhesive in still another layer. Or it could be composed of several drug-in-adhesive layers or combinations of the above layers.

10 The liquid or gel used in the above reservoir type device could be hydrophilic or lipophilic, such as water, alcohols, mineral oils, silicone fluids, various copolymers, such as ethylene vinyl acetate, vinyl acetate or polyvinyl alcohol/polyvinyl pyrrolidone. The reservoir may also include dyes, inert fillers, diluents, antioxidants, penetration enhancers, stabilizers, solubilizing agents and other pharmacologically inactive pharmaceutical agents being well known in the art.

20 The adhesives used are generally of three types, being the rubber type, encompassing inter alia polyisobutylenes, the acrylate type and the silicone type. The adhesives may be chemically modified, may have a wide range of molecular weights. To the adhesive could be added several types of excipients such as fillers, stabilizers, plasticizers, buffering agents, penetration enhancers, penetration retarders, solubilizing agents and other pharmaceutical ingredients being well known in the art.

25 Polymer films which may be used for making the rate-moderating membrane include, without limitation, those comprising low density polyethylene, high density polyethylene, ethyl vinyl acetate copolymers and other suitable polymers.

30 The backing layer serves the purposes of preventing passage of the drug or environmental moisture through the surface of the patch distant from the skin, and also for providing support for the system, where needed. The backing layer may be chosen so that the end product is appealing to the

users, whether children, adults, elderly people or other customer groups. The backing layer is impermeable to the passage of dextromethorphan or inactive ingredients being present in the formulation and can be flexible or nonflexible. Suitable materials include, without limitation, polyester, poly-
5 ethylene terephthalate, some type of nylon, polypropylene, metallized polyester films, polyvinylidene chloride and aluminium foil.

The release liner can be made of the same materials as the backing layer.

As will be clear further below the invention according to the present
10 application encompasses administration of dextromethorphan via all hitherto known types of devices for transdermal administration. Mainly the above categorization will be adhered to in this application. Anyhow this does not exclude that transdermal devices which might fit better according to some other categorization also are included in the present invention.

15 It is well known in the art that the properties of the skin as such influence the penetration of the drug through the skin into the systemic circulation. It could thus be said that the skin controls the drug penetration rate. Anyhow as the skin as such is no part of the present invention the behaviour of the skin in connection with transdermal administration will not be discussed in detail. It is also well accepted in the art that when rate controlling
20 properties are attributed to a transdermal device is meant properties associated with the release rate from the device as such. It is also evident that when a transdermal device is designed to exhibit a certain release performance the properties of the skin need be taken into consideration during the design process.
25

The rate control ability is often a very important feature for a transdermal device in order to deliver the correct drug amount to the patient at the correct time. Thereby maximum efficacy is achieved while side effects are minimized. Many factors influence the rate control ability of a transdermal
30 device. In the below Table 1 the most important such factors are listed and their influence in the respective device type is marked. A plus sign indicates

that the influence is strong. The absence of a plus sign does not exclude that the corresponding has at least some influence.

Table 1

Factor	Type of device			
	Reservoir	Matrix	Drug-in-adhesive	Multilaminate
5 - Polymer type(s)	+	+	+	+
- Modification of the polymer(s)		+	+	+
- Activity, i.e. concentration, of drug e.g. supersaturation	+	+	+	+
10 - Additives in polymer(s)				
Enhancer(s)	+	+	+	+
Cyclodextrine(s)	+	+	+	+
15 Retarder(s)	+	+	+	+
- pH-adjustment	+	+	+	+
- Solubilizer(s)	+	+	+	+
- Emulsifier(s)	+	+	+	+
- Membrane(s)	+			
20 Hydrophilic		+		
Lipophilic	+			
Thickness	+			
Pore size	+			
Density	+			
25 - Chemical stabilizer(s)	+	+	+	+

As a comparably high loading of dextromethorphan is needed for achieving the desirable therapeutic effect the reservoir type device and the multilaminate type device, including several drug-containing layers, are presently considered to be the best modes for carrying out the present transdermal delivery of dextromethorphan.

It is also desirable to include, at least in some device types, one or more transdermal penetration enhancing substance(s) in order to increase the amount of dextromethorphan which may penetrate the skin and which eventually may reach the systemic circulation. Enhancers suitable in the present invention may be categorized in the below groups, although enhancers not belonging to any of these groups are not excluded.

- alcohols, such as short chain alcohols, e.g ethanol and the like, long chain fatty alcohols, e.g. lauryl alcohols, and the like, and polyalcohols, e.g. propylene glycol, glycerine and the like;

- amides, such as amides with long aliphatic chains, or aromatic amides like N,N-diethyl-m-toluamide;

- amino acids;

- azone and azone-like compounds;

- essential oils, i.e. essential oils or constituents thereof, such as l-carvone, l-menthone and the like;

- fatty acids and fatty acid esters, such as oleic acid, lauric acid and the like, further esters of fatty acids, such as isopropyl myristate, and various esters of lauric acid and of oleic acid and the like;

- macrocyclic compounds, such as cyclopentadecanone and cyclodextrins;

- phospholipid and phosphate compounds, such as phospholipids;

- 2-pyrrolidone compounds; and

- miscellaneous compounds, like sulfoxides, such as dimethyl sulfoxides, and fatty acid ethers, such as Laureth-9 and polyoxylaurylether.

Combinations of enhancers from different groups in the above categorization may prove very useful and efficient.

For overviews of enhancers, see further e.g. G.C. Santus et al., "Transdermal enhancer patent literature", Journal of Controlled Release, 1993;25:1-20 and Eric W. Smith et al., "Percutaneous penetration enhancers", CRC Press Inc., 1995.

Detailed description of the invention

The following examples are intended to illustrate but not to limit the scope of the invention, although the embodiments named are of particular interest for our intended purposes.

5 Materials and apparatus used in the examplesMaterials

Dextromethorphan hydrobromide, Roche

Sodium hydroxide, Merck

b-cyclodextrine, Roquette

10 Hydroxypropyl-b-cyclodextrine, Janssen

Isopropyl myristate, Merck

Propylene glycol, Merck

Azone, Discovery Therapeutics Inc.

Ethanol 99.9%, De Danske Spritfabrikker

15 Ethyl acetate, Merck

Disodiumhydrogenphosphate, 2 H₂O, Merck

Polycarbonate membrane 0.2 µm in pore diameter, Whatman

Polycarbonate membrane 0.6 µm in pore diameter, Whatman

Polyester membrane 0.2 µm in pore diameter, Whatman

20 Polyester membrane 0.6 µm in pore diameter, Whatman

Cotran 9702, 3M

Cotran 9711, 3M

Polyester film S 2016, Rexam Release

Polyester film Scotchpak 1220, 3M

25 Polyester film Scotchpak 1109, 3M

MA-24 Medical Grade Adhesive, Adhesives Research Inc.

ETA-2 Medical Grade Adhesive, Adhesives Research Inc.

Eudragit RL100, Röhm GmbH Chemische Fabrik

Eudragit NE 30 D, Röhm GmbH Chemische Fabrik

30 Plastoid E35H, Röhm GmbH Chemische Fabrik

Polyvidone 90, BASF

Durotak 387-2287, National Starch and Chemical B.V.

Apparatus

Franz diffusion cells

Coating equipment: RP Print Coat Instrument LTD., Type KCC 202 Control Coater System with vacuum bed and rods
(100 and 400 μm)

UV-spectrophotometer

Drug Release Apparatus 5, paddle over disk, described in USP 23 p. 1797

HPLC-device: LKB 2248 pump
LKB 2141 variable wavelength monitor
LKB 2221 integrator
LKB 2157 autosampler (20 μl injected)
Precolumn, 4 cm x 4.6 mm i.d., packed with Nucleosil 5 C₁₈
Analytical column, 12 cm x 4.0 mm i.d., packed with Nucleosil 5 C₁₈

The columns were eluted isocratically at ambient temperature with a mobile phase consisting of water-acetonitrile-acetic acid (600:400:1 v/v) with 0.02M potassium nitrate and 0.005M 1-octanesulfonic acid sodium salt. The flow rate was 1.2 ml/min. and the column effluent was monitored at 280 nm.

Example 1

Preparation of dextromethorphan base (in the following called dextromethorphan).

100 g of dextromethorphan hydrobromide was dissolved in 1000 ml of demineralized water. While stirring, the solution was heated to 60°C, until the solution was clear, and approximately 350 ml of sodium hydroxide (1M) was added (the addition of sodium hydroxide (1M) was stopped when precipitation no longer occurred). The precipitated mixture was refrigerated for at least 4 hours. The mixture was vacuum filtered and dried in a drying oven at 25°C. The resulting dextromethorphan passed the test of USP 23, p. 481.

Example 2

Analysis of the receptor solutions described in Examples 3 and 4.

Quantitative determination of dextromethorphan in the receptor solution samples from the skin permeation studies in Example 3 and from the membrane permeation studies in Example 4 was done by the HPLC method described under Apparatus.

Example 3

In vitro skin permeation studies from solutions of dextromethorphan.

Solution 1

A saturated dextromethorphan solution in demineralized water.

Solution 2

A saturated dextromethorphan solution in demineralized water containing 10 mg/ml of b-cyclodextrine.

Solution 3

A saturated dextromethorphan solution in demineralized water containing 10 mg/ml of hydroxypropyl-b-cyclodextrine.

Solution 4

50 mg dextromethorphan was dissolved in 5 ml isopropyl myristate.

Solution 5

50 mg dextromethorphan was dissolved in 5 ml propylene glycol.

Solution 6

50 mg dextromethorphan was dissolved in 5 ml propylene glycol containing 50 mg/ml of azone.

Solution 7

250 mg dextromethorphan was dissolved in 5 ml ethanol.

Solution 8

150 mg dextromethorphan was dissolved in 5 ml ethyl acetate.

In vitro permeation of dextromethorphan from the solutions 1, 2, 3, 4, 5, 6, 7 and 8 through dermatomed pig skin was investigated in Franz diffusion Cells.

Skin pieces with a thickness of approximately 765 μm were dermatomed from full thickness pig skin and mounted in glass diffusion cells with an available diffusion area of 1.8 cm^2 . Pig skin is a fully accepted model for human skin. The solutions were applied separately on the skin surfaces and the dermal sides were all exposed to 12.1 ml receptor solution consisting of a 0.05M phosphate buffer solution of pH 7.4 equilibrated to $37 \pm 1^\circ\text{C}$.

Permeation of dextromethorphan was followed by removing samples periodically and measuring the concentration by the HPLC method according to Example 2. The cumulative amount of dextromethorphan appearing in the receptor solution versus time is shown in FIG. 2. An increase in the permeated amount of dextromethorphan is seen in the following order: Water, isopropyl myristate, propylene glycol, propylene glycol containing 5 % azone, water containing 1% hydroxpropyl-b-cyclodextrine, water containing 1% b-cyclodextrine, ethanol and ethyl acetate used as solvents. The maximal observed flux of dextromethorphan is 21 $\mu\text{g}/\text{cm}^2/\text{h}$ and the range is from approximately 0.5 to 21 $\mu\text{g}/\text{cm}^2/\text{h}$.

The results show that dependent on the used solvent it is possible to optimize the flux of dextromethorphan through the skin. Both by using ethanol and ethyl acetate and by addition of cyclodextrines, a remarkable increase in the fluxes is seen.

Example 4

In vitro permeation studies across artificial membranes from solutions of dextromethorphan, imitating the reservoir type transdermal device.

Solution 9

50 mg dextromethorphan was dissolved in 5 ml propylene glycol.

Solution 10

50 mg dextromethorphan was dissolved in 5 ml ethanol.

In vitro permeation of dextromethorphan from the solutions 9 and 10 across 6 different types of artificial membranes was investigated in Franz diffusion cells.

Artificial membranes of the following types were studied: Whatman 0.2 μm PC (polycarbonate), Whatman 0.6 μm PC (polycarbonate), Whatman 0.2 μm PET (polyester), Whatman 0.6 μm PET (polyester), Cotran 9702 (ethylene vinyl acetate film) and Cotran 9711 (microporous polyethylene film). The membranes were mounted in glass diffusion cells with an available diffusion area of 1.8 cm^2 . Solution 9 was applied on the surface of all the above membranes while solution 10 only was applied on Cotran 9702 and Cotran 9711. The opposite sides of the membranes were all exposed to 12.1 ml receptor solution consisting of a 0.05M phosphate buffer solution of pH 7.4 equilibrated to $37 \pm 1^\circ\text{C}$.

Permeation of dextromethorphan was followed by removing samples periodically and measuring the concentration by the HPLC method according to Example 2. The cumulative amount of dextromethorphan appearing in the receptor solution versus time is shown in FIG. 3. An increase in the permeated amount of dextromethorphan was seen in the following order of used membranes: Cotran 9702, Whatman 0.2 μm PC, Whatman 0.2 μm PET, Cotran 9711, Whatman 0.6 μm PC and Whatman 0.6 μm PET. The results show that it is possible to control the release rate of dextromethorphan through different membrane type. On this basis it is easy to produce a reservoir type device by sealing the membrane to a backing layer being impermeable to dextromethorphan and other components of the formulation.

Example 5

Transdermal drug delivery systems with dextromethorphan or dextromethorphan hydrobromide as the active substances

System 1 (drug-in-adhesive type, acrylate)

2.5 g dextromethorphan was suspended in 20 g ETA-2 Medical Grade Adhesive to give the drug gel. The drug gel was solvent cast onto a polyester film, S 2016, by means of the coating equipment (wet layer = 400 μm). After drying at 80°C for 10 minutes, a polyester film, Scotchpak 1109, was laminated onto the dried drug gel. The resulting sheet was die-cut into patches

which was kept at room temperature until use. The concentration of dextromethorphan was approximately 1,5 mg/cm².

System 2 (drug-in-adhesive type, acrylate)

5 5 g dextromethorphan was dissolved in 10 ml ethanol. The solution was added to 15 g Durotak 387-2287 to give the drug gel. The drug gel was solvent cast onto a polyester film, S 2016, by means of the coating equipment (wet layer = 400 µm). After drying at 80°C for 10 minutes, a polyester film, Scotchpak 1220, was laminated onto the dried drug gel. The resulting sheet was die-cut into patches which were kept at room temperature until
10 use. The concentration of dextromethorphan was approximately 2 mg/cm².

System 3 (multi-laminate type, waterbased acrylate)

2.4 g dextromethorphan hydrobromide was dispersed in a mixture of 3 g Eudragit NE 30 D and 45 g Plastoid E35H to give the drug gel. The drug gel was solvent cast onto a polyester film, S 2016, by means of the coating
15 equipment (wet layer = 400 µm). After drying at 80°C for 10 minutes, an adhesive layer consisting of Plastoid E35H (wet layer = 100 µm) coated on a polyester film, S 2016, was laminated onto the dried drug gel. The polyester film, S 2016, in contact with the drug gel was removed, and Scotchpak 1109 was laminated onto the drug gel as the backing. The resulting sheet was die-
20 cut into patches which were kept at room temperature until use. The concentration of dextromethorphan was approximately 1 mg/cm².

System 4 (drug-in-adhesive type, acrylate)

2.5 g dextromethorphan was suspended in 20 g Durotak 387-2287 to give the drug gel. The drug gel was solvent cast onto a polyester film, S 2016,
25 by means of the coating equipment (wet layer = 400 µm). After drying at 80°C for 10 minutes, a polyester film, Scotchpak 1109, was laminated onto the dried drug gel. The resulting sheet was die-cut into patches which were kept at room temperature until use. The concentration of dextromethorphan was approximately 2 mg/cm².

System 5 (drug-in-adhesive type, waterbased acrylate)

2.4 g dextromethorphan hydrobromide was dispersed in a mixture of 3 g Eudragit NE 30 D and 45 g Plastoid E35H to give the drug gel. The drug gel was solvent cast onto a polyester film, S 2016, by means of the coating equipment (wet layer = 400 μm). After drying at 80°C for 10 minutes, a polyester film, Scotchpak 1109, was laminated onto the dried drug gel. The resulting sheet was die-cut into patches which were kept at room temperature until use. The concentration of dextromethorphan was approximately 1 mg/cm².

System 6 (multi-laminate type, acrylate)

2.5 g dextromethorphan was dissolved in 10 ml ethanol. The solution was added to a mixture of 12.8 g Eudragit gel (50 % Eudragit RL 100 swelled in ethanol), 12.8 g PVP gel (20 % Polyvidone 90 swelled in ethanol) and 4 g propylene glycol to give the drug gel. The drug gel was solvent cast onto a polyester film, S 2016, by means of the coating equipment (wet layer = 400 μm). After drying at 80°C for 10 minutes, an adhesive layer consisting of Plastoid E35H (wet layer = 100 μm) coated on a polyester film, S 2016, was laminated onto the dried drug gel. The polyester film, S 2016, in contact with the drug gel was removed, and Scotchpak 1109 was laminated onto the drug gel as the backing. The resulting sheet was die-cut into patches which were kept at room temperature until use. The concentration of dextromethorphan was approximately 0.5 mg/cm².

System 7 (drug-in-adhesive type, polyisobutylene)

2.5 g dextromethorphan was suspended in 10 g MA-24 Medical Grade Adhesive to give the drug gel. The drug gel was solvent cast onto a polyester film, S 2016, by means of the coating equipment (wet layer = 400 μm). After drying at 80°C for 10 minutes, a polyester film, Scotchpak 1220, was laminated onto the dried drug gel. The resulting sheet was die-cut into patches which were kept at room temperature until use. The concentration of dextromethorphan was approximately 2 mg/cm².

In vitro release studies according to Example 6 were carried out on the systems 1, 2, 3, 4, 5, 6 and 7 described above. The results of these studies are shown graphically in FIG. 4.

The results show that different release profiles can be achieved from
5 different types of devices.

Example 6

In vitro release studies of the transdermal drug delivery systems according to Example 5.

The apparatus used was Apparatus 5, paddle over disk, described under
10 Apparatus. Patches of 7.1 cm² were applied to the disk assembly, using a suitable adhesive, with the release surface facing up. The dissolution medium used was 600 ml of 0.05M phosphate buffer pH 7.4 equilibrated to 32±0.5°C. Samples were withdrawn at 1, 2, 4, 8 and 24 hours, respectively. The amount of dextromethorphan in the samples was determined by UV-
15 spectrophotometry at 280 nm and the concentration of the respective systems was expressed in mg dextromethorphan per cm².

A reservoir type device may be manufactured by heat sealing a membrane such as described in the above Example 4 to a backing containing the drug in a suitable vehicle.

20 A iontophoretic type device may be manufactured essentially according to embodiments disclosed in e.g. Parminder Singh et al, "Iontophoresis in Drug Delivery: Basic Principles and Applications", Critical Reviews in Therapeutic Drug Carrier Systems, 1994; 11 (2&3):161-213. The administration of dextromethorphan is not disclosed in this reference. Anyhow it lies within
25 the present invention to modify, using the disclosure in the present application, the embodiments according to said reference to become suitable for the administration of dextromethorphan.

The above examples show that it is possible to administer dextromethorphan and to control its release rate using all known types of devices for transdermal drug administration.
30

Some prodrug type derivatives of dextromethorphan can be used according to the present invention for obtaining the desirable antitussive effect. Such derivatives may include other ethers in the 3-position than the methoxy-group. By modification in the 3-position compounds with favourable permeation rates through human and animal skin may be obtained. Upon permeation of stratum corneum dextromethorphan or dextrorphan may be generated through metabolic reactions. Other salts than the hydrobromide could be used as it is known that more lipophilic anions than bromide may generate ion-pairs with more favourable skin permeation rates.

It is evident that the above mentioned Examples may be modified to encompass also metabolites, different salts and prodrugs of dextromethorphan.

Various carriers and vehicles for dextromethorphan may be used in the transdermal administration. One such carrier is cyclodextrin, especially β -cyclodextrin. Dextromethorphan can be bound in the cavities of cyclodextrins to form so called inclusion complexes. Binding dextromethorphan to a cyclodextrin leads either to increased delivery rate or to decreased delivery rate depending on the dextromethorphan-cyclodextrine ratio.

It is within the present invention to add to the transdermal device substances being fragrances or other substances with agreeable smell in order to give the device a smell appealing to the user.

As the period of time from the first application of a transdermal device according to the present invention until a therapeutically effective serum level of dextromethorphan is achieved is in the order of up to 3 hours the complementary and concomitant use of another administration form may be of value. Oral, sublingual, buccal, nasal, pulmonary and rectal, and possibly other transmucosal, administration of dextromethorphan results in that the drug reaches the system more rapidly than through the transdermal route. As mentioned above said non-transdermal administration forms have the disadvantage of a lower bioavailability than the transdermal form of admini-

stration. Anyhow this disadvantage, and problems related thereto, may be temporarily tolerated if an antitussive effect is desirable in the period of time before the therapeutic effect is achieved from the transdermal device.

One suitable use of the mentioned forms of administration is to administer dextromethorphan through the oral, sublingual, buccal, nasal, pulmonary or rectal, or possibly other transmucosal routes approximately at the same time as the first transdermal device is applied. Thereafter new transdermal devices are applied to ensure the correct plasma level without further administration through the oral, sublingual, buccal, nasal, pulmonary and rectal, or possibly other transmucosal, route. The above concomitant use of different administration forms is especially useful in certain situations, such as, but not exclusively, some time prior to oral presentations, attendance to conferences and visits to theatres, concerts and church. It is thus feasible to market set of formulations including devices for transdermal administration as well as devices or formulations for oral, sublingual, buccal, nasal, rectal, pulmonary and rectal, and possibly other transmucosal, administration of dextromethorphan.

Another envisageable concomitant use according to the present invention is to apply a second transdermal device while a priorly applied first transdermal device is still adhered to the patient's skin while still delivering some amount of the drug. The utility behind this use is as follows. Suppose that the transdermal devices used deliver the drug during 36 hours. The first evening one such device is applied. The following evening the device still delivers the drug, though usually with a lower flux rate than earlier. If now this second evening a second transdermal device is applied while the first one is left on the skin the fluxes from the first and second device will add to a useful flux as the flux from the first device successively decreases while the drug from the second device only reaches the systemic circulation after some hours. By using transdermal devices in this way a more stable therapeutically effective plasma level of the drug during an extended period of time is achieved than if for example are used devices delivering for 24 hours

and being replaced every 24 hours. Also other useful combinations of concomitantly used transdermal devices are envisageable.

As it might be advantageous that the cough now and then should be allowed to occur it might be desirable not to treat or prevent cough during too long continuous periods of time. It is within the present invention to administer dextromethorphan in such a way that a therapeutically effective systemic level of dextromethorphan prevails mainly during those periods of time during day and night when it is desirable that cough should be treated or prevented, and, consequently, in such a way that a less than therapeutically effective systemic level of dextromethorphan prevails mainly during those periods of time during day and night when it is not desirable that cough should be treated or prevented. The above object is achievable by applying the transdermal device at the appropriate time during day or night in combination with designing the device with the appropriate release profile.

15 Dosage

The maximal dose of dextromethorphan to be given perorally according to Martindale, "The Extra Pharmacopoeia", London, 1993 is for adults 10 to 30 mg every 4 to 8 hours up to a maximum of 120 mg in every 24 hours. Children aged 6 to 12 years may be given 5 to 15 mg perorally every 4 to 8 hours to a maximum in 24 hours and children aged 1 to 6 years 2.5 to 7.5 mg every 4 to 8 hours to a maximum of 30 mg in 24 hours. Similar peroral dosages are recommended in "Handbook of Non-prescription Drugs", 10th ed., American Pharmaceutical Association, The National Professional Society of Pharmacists, Washington D.C., 1993.

25 The average hourly flux to be achieved from a transdermal formulation can be calculated from an average oral dosage of 60 mg in every 24 hours and a bioavailability of 25 % in oral delivery. Assuming 100 % bioavailability in using the transdermal route the average 24 hours transdermal dose should be $0,25 \times 60 \text{ mg} = 15 \text{ mg}$, which corresponds to an hourly flux of
30 $(15 \times 1000)/(30 \times 24) = 21 \text{ mg/cm}^2/\text{hour}$ from a trans-dermal device with an area of 30 cm^2 . Recalculating to transdermal delivery corresponding to the

maximum oral dosage of 120 mg in every 24 hours results in an hourly flux of 42 mg/cm²/hour from a transdermal device with an area of 30 cm².

The area of a transdermal device being convenient for a patient to wear is in the range from 5 to 50 cm². The corresponding patch loading should be at least 0,3 - 1,5 mg/cm² for a transdermal device with an area of 30 cm². As the drug content of a transdermal device is never completely depleted during its application to a patient a higher loading than above must be anticipated, preferably 1 - 3 mg/cm². The above indicated loadings in mg/cm² are to be considered as average loadings for an average size device. It is known that the driving force for the release of a drug from a transdermal device is related to the drug concentration, i.e. number of mg of drug/cm³. Therefore the above indicated loadings in mg/cm² are to be adjusted according to the actual areal size and thickness of the device in order to arrive at the desirable therapeutic effect.

Loadings for different sizes and types of devices for transdermal administration, taking into account different age groups and types of patients, range from about 0.1 mg/cm² to about 10 mg/cm² of dextromethorphan. The hourly flux rate of dextromethorphan ranges from about 1 mg/cm²/hour to about 100 mg/cm²/hour. The effective transdermally delivered amount of dextromethorphan is from about 0.05 mg/kg bodyweight to about 5 mg/kg bodyweight.

It should also be contemplated that a device for transdermal delivery during 8 - 12 hours would be clinically more relevant than a device for delivery during 24 hours. Such a device with limited release duration may be used for the periods with worst cough during daytime and during the night allowing the patient to cough the bronchi clean therebetween.

The mentioned device may either be taken off from the skin after 8 - 12 hours in order to stop further delivery, or be designed in such a way that its delivery drops to negligible or non-pharmacological levels after 8 - 12 hours. In this latter case the device may remain on the skin after 8 - 12 hours without the patient risking further delivery thereafter which facilitates the

patient's handling of the device. Such devices are known per se, see eg US 4,915,950 (MIRANDA ET AL.) - although not for delivery of dextromethorphan.

5 When dextromethorphan is administered in a transdermal device the latter should preferably be occlusive, which means that the device does not permit water to migrate outwardly from the patient. Thereby the hydration of the skin is increased which favours the penetration of dextromethorphan through the skin.

CLAIMS

1. Device for transdermal administration, characterized in that it administers dextromethorphan, (+)-3-methoxy-17-methyl-9a,13a,14a-morphinan, optionally encompassing salts, prodrugs and metabolites thereof, and optionally together with pharmaceutically acceptable carrier(s), to a human being or an animal in order to achieve an antitussive effect.
2. Device for transdermal administration according to claim 1, characterized in that it is of the reservoir type, the matrix type, the drug-in-adhesive type, the multi-laminate type and/or the iontophoretic type or combinations thereof, preferably of the reservoir type or the multi-laminate type or combinations of these two types.
3. Device for transdermal administration according to anyone of the preceding claims, characterized in that it has a loading of dextromethorphan from about 0.1 mg/cm² to about 10 mg/cm².
4. Device for transdermal administration according to anyone of the preceding claims, characterized in that it has an hourly flux rate of dextromethorphan from about 1 mg/cm²/hour to about 100 mg/cm²/hour.
5. Device for transdermal administration according to anyone of the preceding claims, characterized in that it delivers dextromethorphan for a predefined period of time, preferably 8, 12 or 24 hours.
6. Device according to anyone of the preceding claims, characterized in that dextromethorphan is present in a complex with cyclodextrin, preferably b-cyclodextrin.

7. Device according to anyone of the preceding claims, c h a r a c -
t e r i z e d in that it has a release profile being such that it, when applied on
the skin at the appropriate time during day or night, administers dex-
tromethorphan in such a way that a therapeutically effective systemic level
5 of dextromethorphan prevails mainly during such periods of time during day
and night when an antitussive effect is most desirable.

8. Device according to anyone of the preceding claims, c h a r a c -
t e r i z e d in that it further comprises a fragrance or other substance with
agreeable smell.

10

9. Device according to anyone of the preceding claims, c h a r a c -
t e r i z e d in that it further comprises a substance enhancing transdermal
penetration.

15

10. Device according to anyone of the preceding claims, c h a r a c -
t e r i z e d in that it is occlusive.

11. Set of formulations of dextromethorphan, (+)-3-methoxy-17-
methyl-9a,13a,14a-morphinan, optionally encompassing salts, prodrugs and
20 metabolites thereof, and optionally together with pharmaceutically accept-
able carrier(s), c h a r a c t e r i z e d in that it comprises at least one device
according to anyone of the preceding claims and at least one formulation for
oral, sublingual, buccal, nasal, pulmonary, rectal and/or other transmucosal
administration.

25

12. Use of an antitussive compound comprising dextromethorphan,
(+)-3-methoxy-17-methyl-9a,13a,14a-morphinan, optionally encompassing
salts, prodrugs and metabolites thereof, and optionally together with phar-
maceutically acceptable carrier(s), for the manufacture of a composition to
30 be administered transdermally for achieving an antitussive effect.

13. Use according to claim 12, characterized in that the transdermal delivery is carried out using a device for transdermal delivery, such device especially being of the reservoir type, the matrix type, the drug-in-adhesive type, the multi-laminate type and/or the iontophoretic type or combinations thereof, preferably of the reservoir type or the multi-laminate type or combinations of these two types.

14. Use according to claim 13, characterized in that more than one device transdermal device is used at a time.

10

15. Method for achieving an antitussive effect in a living body by transdermal administration of a compound comprising dextromethorphan, (+)-3-methoxy-17-methyl-9a,13a,14a-morphinan, optionally encompassing salts, prodrugs and metabolites thereof, and optionally together with pharmaceutically acceptable carrier(s).

16. Method according to claim 15 wherein the treatment is achieved through systemic effect of the transdermally administered compound.

17. Method according to anyone of claims 15 or 16 wherein the transdermal administration is carried out using a device for transdermal delivery, such device especially being of the reservoir type, the matrix type, the drug-in-adhesive type, the multi-laminate type and/or the iontophoretic type or combinations thereof, preferably of the reservoir type or the multi-laminate type or combinations of these two types.

18. Method according to anyone of claims 15 - 17 wherein more than one device for transdermal delivery is used at a time.

19. Method according to anyone of claims 15 - 18 wherein the effective transdermally delivered amount of dextromethorphan is from about

0.05 mg/kg bodyweight to about 5 mg/kg bodyweight during a predefined period of time, preferably 8, 12 or 24 hours.

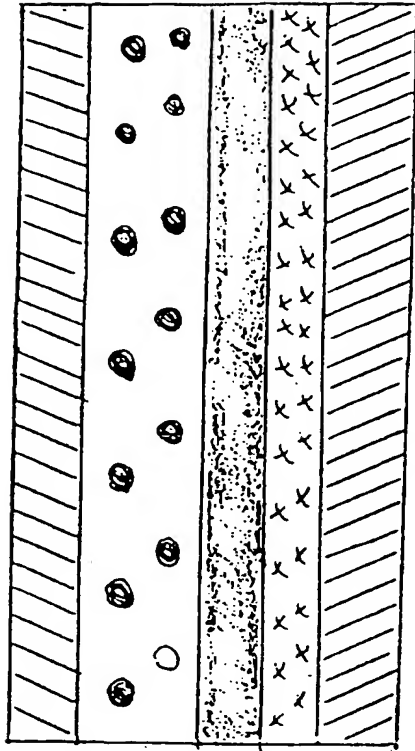
20. Method for achieving an antitussive effect in a living body by
5 transdermal administration of a compound comprising dextromethorphan, (+)-3-methoxy-17-methyl-9a,13a,14a-morphinan, optionally encompassing salts, prodrugs and metabolites thereof, and optionally together with pharmaceutically acceptable carrier(s) in combination with oral, sublingual, buccal, nasal, pulmonary, rectal and/or other transmucosal administration of a
10 compound comprising dextromethorphan, (+)-3-methoxy-17-methyl-9a,13a,14a-morphinan, optionally encompassing salts, prodrugs and metabolites thereof, and optionally together with pharmaceutically acceptable carrier(s).

15. 21. Method according to anyone of the claims 15 -20, c h a r a c -
t e r i z e d in that dextromethorphan is administered in such a way that a therapeutically effective systemic level of dextromethorphan prevails mainly during those periods of time during day and night when an antitussive effect is most desirable.

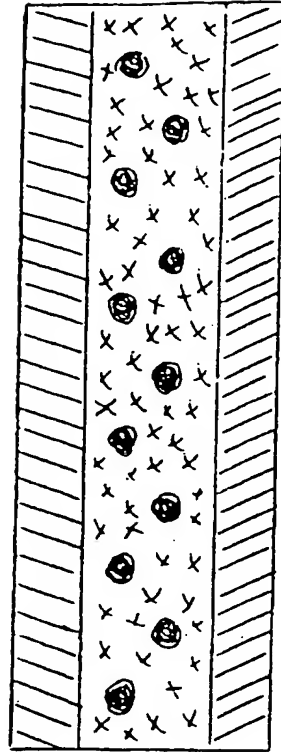
20

22. Method according to anyone of the claims 15 - 21, c h a r a c -
t e r i z e d in that dextromethorphan is administered in such a way that a less than therapeutically effective systemic level of dextromethorphan prevails mainly during those periods of time during day and night when an
25 antitussive effect is less desirable.

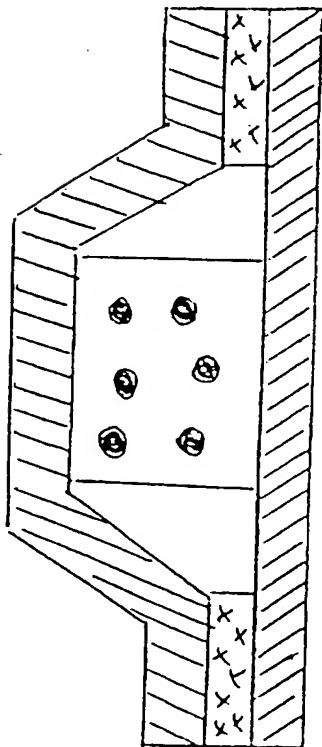
1/4



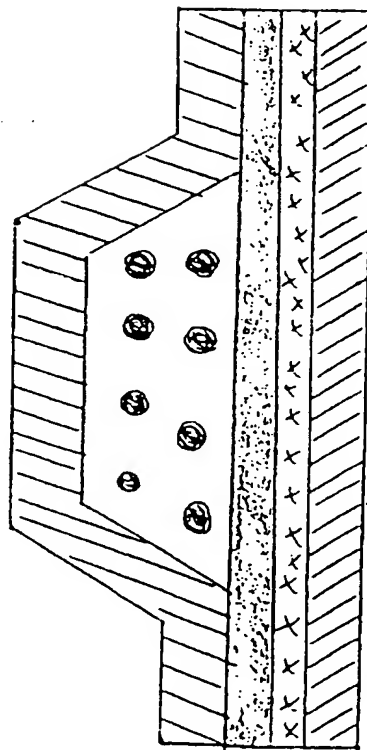
Multi-laminate Fig. 1 B



Drug-in-adhesive Fig. 1 D



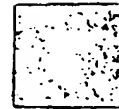
Matrix Fig. 1 A



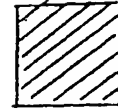
Reservoir Fig. 1 C



Backing



Membrane



Liner/skin

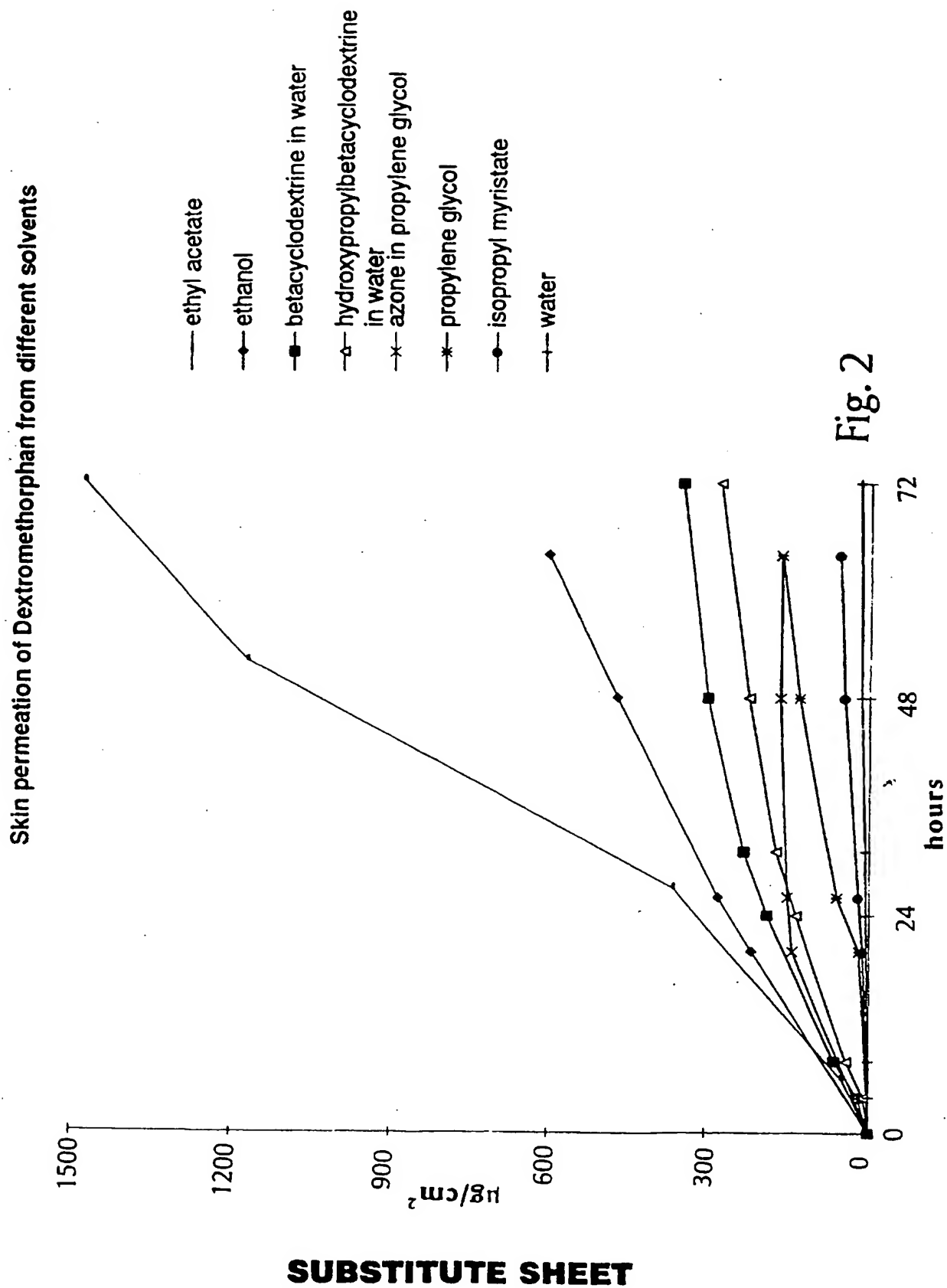


Drug



Adhesive

2/4



3/4

Membrane permeation of Dextromethorphan

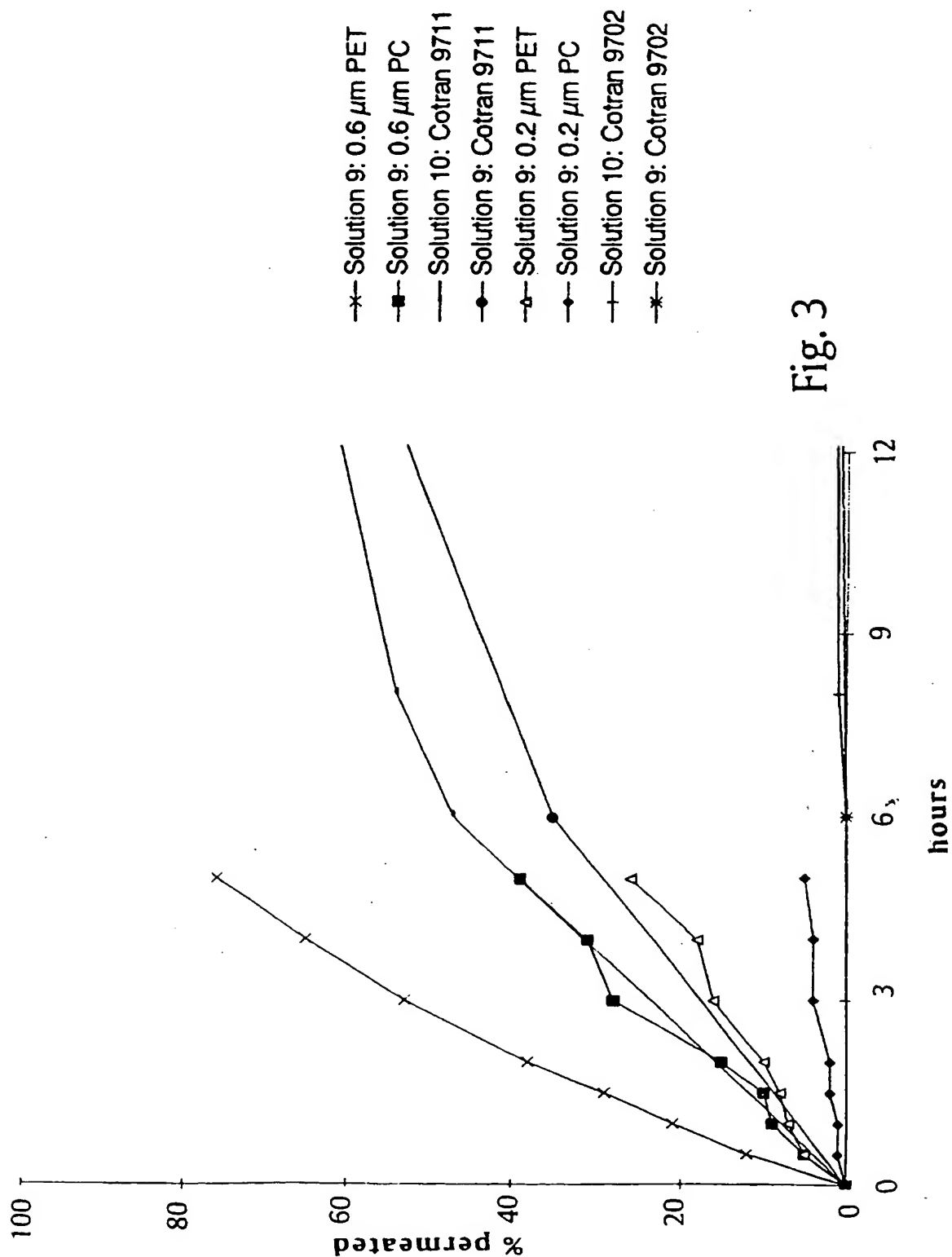


Fig. 3

SUBSTITUTE SHEET

4/4

Release of Dextromethorphan from transdermal systems

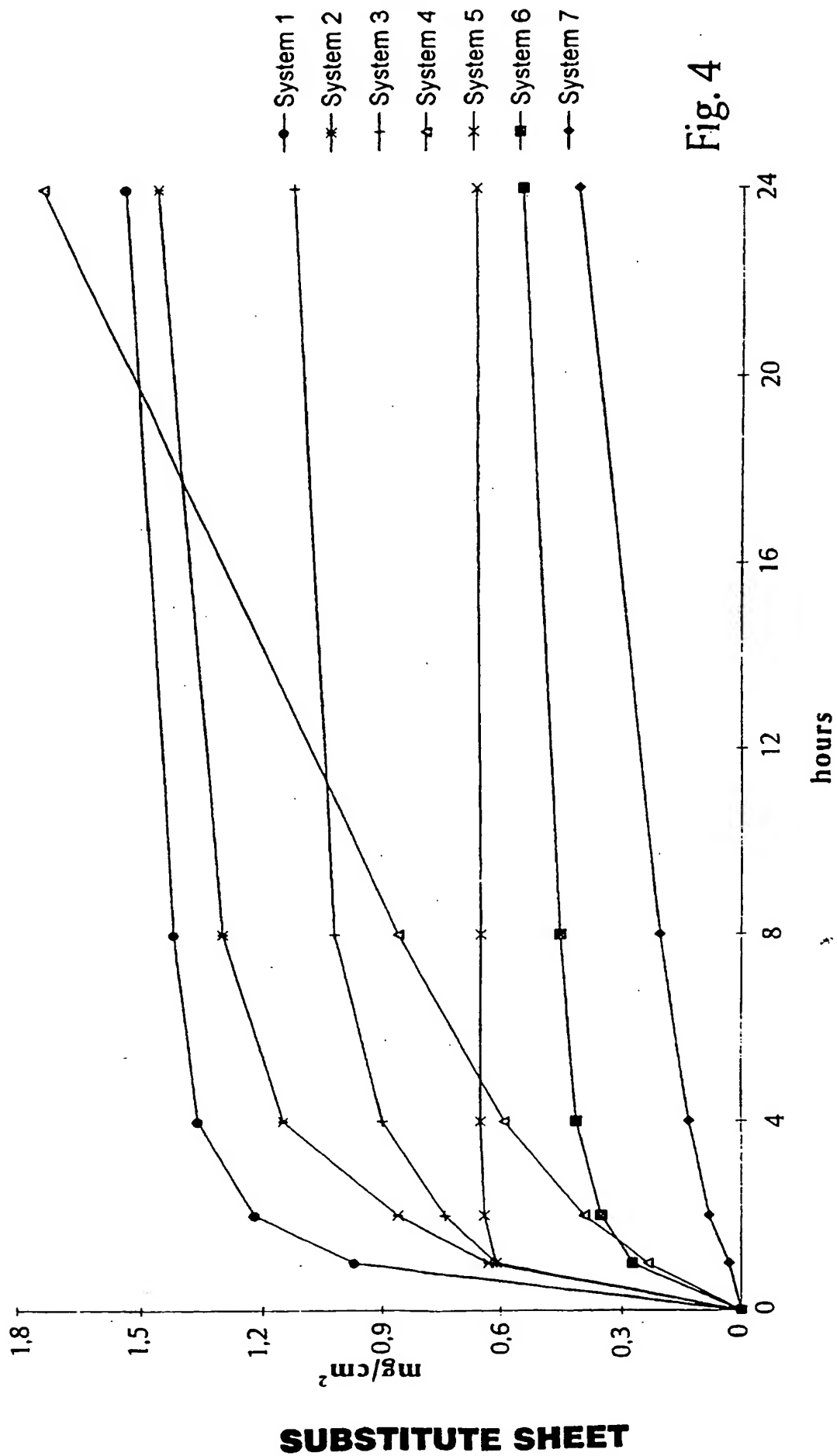


Fig. 4

SUBSTITUTE SHEET

INTERNATIONAL SEARCH REPORT

International application No.:

PCT/SE 97/00484

A. CLASSIFICATION OF SUBJECT MATTER

IPC6: A61K 9/70, A61K 31/485

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC6: A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAPLUS, USFULLTEXT, WPI, WPIL

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 9115261 A1 (MEDTRONIC, INC.), 17 October 1991 (17.10.91), page 1, line 10 - line 14; page 4, line 32 - line 33	1-10, 12-19, 21-22 PARTLY
Y	--	11, 20, 21-22 PARTLY
X	US 5260066 A5 (LOUIS L. WOOD ET AL), 9 November 1993 (09.11.93), claims	1-10, 12-19, 21-22 PARTLY
Y	--	11, 20, 21-22 PARTLY
Y	WO 9214466 A1 (SMITHKLINE BEECHAM PLC), 3 Sept 1992 (03.09.92), claims	11, 20, 21-22 PARTLY
	--	

☒ Further documents are listed in the continuation of Box C.

☒ See patent family annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

30 June 1997

Date of mailing of the international search report

05 -07- 1997

Name and mailing address of the ISA/
Swedish Patent Office
Box 5055, S-102 42 STOCKHOLM
Facsimile No. +46 8 666 02 86

Authorized officer

Anneli Jönsson
Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 97/00484

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 9207559 A1 (THE PROCTER & GAMBLE COMPANY), 14 May 1992 (14.05.92), claims --	11,20, 21-22 PARTLY
A	WO 8807871 A1 (WARNER-LAMBERT COMPANY), 20 October 1988 (20.10.88) -- -----	1-10,12-14

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 97/00484

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 15-22
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Claims 15-22 are directed to methods of treatment of the human or animal body by surgery or by therapy/diagnostic methods practised on the human or animal body/Rule 39.1(iv). Nevertheless, a search has been executed for these claims. The search has been based on the alleged effects of the compounds/compositions.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. Claims 1-10, 12-19 and 21-22 (partly) device, use and method of transdermal administration of dextromethorphan as an antitussive agent.
 2. Claims 11, 20 and 21-22 (partly) set of formulations for administration of dextromethorphan, i.e. combination of different ways of administrations.
1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
 2. ☒ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
 3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
 4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT
Information on patent family members

03/06/97

International application No.

PCT/SE 97/00484

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9115261 A1	17/10/91	AT 142116 T AU 649439 B CA 2079007 A DE 69121885 D,T EP 0522062 A,B US 5213568 A	15/09/96 26/05/94 01/10/91 20/03/97 13/01/93 25/05/93
US 5260066 AS	09/11/93	US 5288503 A	22/02/94
WO 9214466 A1	03/09/92	AU 1227292 A	15/09/92
WO 9207559 A1	14/05/92	AU 663857 B AU 8915891 A CA 2129676 A CZ 9300795 A EP 0555411 A FI 931934 A HU 64216 A JP 6502428 T PL 168968 B SK 42593 A US 5196436 A	26/10/95 26/05/92 14/05/92 19/01/94 18/08/93 31/05/93 28/12/93 17/03/94 31/05/96 09/09/93 23/03/93
WO 8807871 A1	20/10/88	DE 3887156 D,T EP 0382716 A,B SE 0382716 T3 JP 2502994 T US 4783450 A	28/04/94 22/08/90 20/09/90 08/11/88